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Amendments to the Claims

This listing of claims will replace all prior versions, and listings, of claims in the application:

Listing of Claims

1-6 (Canceled)

7. (Currently Amended) A method of specifically inhibiting fusion of a CD4+ cell susceptible to infection by a macrophage-tropic primary isolate of HIV-1 to an envelope of a macrophage-tropic primary isolate of HIV-1 to a CD4+ cell susceptible to infection by a macrophage-tropic primary isolate of HIV-1 which comprises contacting the CD4+ cell with an agent which is (1) capable of inhibiting fusion of HeLa-env<sub>JR-FL</sub> to a PM1 cell, but (2) not capable of inhibiting fusion of HeLa-env<sub>LAI</sub> to a HeLa-CD4+ cell, so as to thereby inhibit the wherein the agent inhibits fusion of the CD4+ cell to the envelope of the macrophage-tropic primary isolate of HIV-1 to the CD4+ cell.

8. (Previously Presented) The method of claim 7, wherein the agent is determined to be capable of inhibiting fusion of a macrophage-tropic primary isolate of HIV-1 to a CD4+ cell but not capable of inhibiting fusion of a T cell tropic isolate of HIV-1 to a CD4+ cell using a method which comprises:

(a) contacting (i) a PM1 cell, which is labeled with a first dye, with (ii) HeLa-env<sub>JR-FL</sub>, which is labeled with a second dye, in the presence of an excess of the agent under conditions which would normally permit the fusion of the PM1 cell to the HeLa-env<sub>JR-FL</sub> in the absence of the agent, the first and second dyes being

selected so as to allow resonance energy transfer between the dyes;

- (b) exposing the product of step (a) to conditions which would result in resonance energy transfer if fusion has occurred; and
- (c) determining whether there is a reduction of resonance energy transfer, when compared with the resonance energy transfer in the absence of the agent;
- (d) contacting (i) a HeLa-CD4+ cell, which is labeled with a first dye, with (ii) HeLa-env<sub>LAI</sub> which is labeled with a second dye, in the presence of an excess of the agent under conditions which would normally permit the fusion of HeLa-CD4+ to the HeLa-env<sub>LAI</sub> in the absence of the agent, the first and second dyes being selected so as to allow resonance energy transfer between the dyes;
- (e) exposing the product of step (d) to conditions that would result in resonance energy transfer if fusion has occurred;
- (f) determining whether there is a reduction of resonance energy transfer, when compared with the resonance energy transfer in the absence of the agent; and
- (g) comparing the determination made in step (c) with the determination made in step (f), wherein a decrease in transfer in step (c) but not in step (f) indicates that the agent is capable of specifically inhibiting fusion of the macrophage-tropic primary isolate of HIV-1 to the CD4+ cell, but not capable of specifically inhibiting the fusion of a T cell-tropic isolate of HIV-1 to the CD4+ cell.

9. (Previously Presented) The method of claim 7, wherein the agent is an antibody.

10-12. (Canceled)

13. (Previously Presented) The method of claim 7, wherein the agent is capable of inhibiting fusion of a macrophage-tropic primary isolate of HIV-1 to a CD4+ cell but not capable of inhibiting fusion of a T cell-tropic isolate of HIV-1 to a CD4+ cell in a method which comprises:

- (a) contacting (i) a PM1 cell, which is labeled with a first dye, with (ii) HeLa-env<sub>JR-FL</sub>, which is labeled with a second dye, in the presence of an excess of the agent under conditions which would normally permit the fusion of the PM1 cell to the HeLa-env<sub>JR-FL</sub> in the absence of the agent, the first and second dyes being selected so as to allow resonance energy transfer between the dyes;
- (b) exposing the product of step (a) to conditions which would result in resonance energy transfer if fusion has occurred; and
- (c) determining whether there is a reduction of resonance energy transfer, when compared with the resonance energy transfer in the absence of the agent;
- (d) contacting (i) a HeLa-CD4+ cell, which is labeled with a first dye, with (ii) HeLa-env<sub>LAI</sub>, which is labeled with a second dye, in the presence of an excess of the agent under conditions which would normally permit the fusion of HeLa-CD4+ to the HeLa-env<sub>LAI</sub> in the absence of the agent, the first and second dyes being selected so as to allow resonance energy transfer between the dyes;
- (e) exposing the product of step (d) to conditions that would result in resonance energy transfer if fusion has occurred;
- (f) determining whether there is a reduction in resonance energy transfer, when compared with the resonance

(g) energy transfer in the absence of the agent; and  
comparing the determination made in step (c) with the  
determination made in step (f), wherein a decrease in  
transfer in step (c) but not in step (f) indicates  
that the agent is capable of specifically inhibiting  
fusion of the macrophage-tropic primary isolate of  
HIV-1 to the CD4+ cell, but not capable of  
specifically inhibiting the fusion of a T cell-tropic  
isolate of HIV-1 to the CD4+ cell.

14. (Previously Presented) The method of claim 7, wherein the  
agent is a protein moiety.

15. (Previously Presented) The method of claim 14, wherein the  
protein moiety is an antibody.

16. (Previously Presented) The method of claim 15, wherein the  
antibody is an antibody is a monoclonal antibody.

17. (Previously Presented) The method of claim 15, wherein the  
antibody is a wholly synthetic antibody or a chimeric  
antibody.

18. (Previously Presented) The method of any of claims 15-17,  
wherein the antibody is an antigen-binding fragment of an  
antibody.

19. (Previously Presented) The method of claim 14, wherein the  
protein moiety is a  $\beta$ -chemokine.

20. (Currently Amended) A method of specifically inhibiting  
fusion of a CD4+ cell susceptible to infection by a  
macrophage-tropic primary isolate of HIV-1 to an envelope  
of a macrophage-tropic primary isolate of HIV-1 to a CD4+

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~~cell susceptible to infection by a macrophage-tropic primary isolate of HIV-1 which comprises contacting the CD4+ cell with a protein moiety which is (1) capable of inhibiting fusion of HeLa-env<sub>JR-FL</sub> to a PM1 cell, but (2) not capable of inhibiting fusion of HeLa-env<sub>LA1</sub> to a HeLa-CD4+ cell, so as to thereby inhibit the wherein the protein moiety inhibits fusion of the CD4+ cell to the envelope of the macrophage-tropic primary isolate of HIV-1 to the CD4+ cell.~~

21. (Previously Presented) The method of claim 20, wherein the protein moiety is an antibody.
22. (Previously Presented) The method of claim 21, wherein the antibody is a monoclonal antibody.
23. (Previously Presented) The method of claim 21, wherein the antibody is a wholly synthetic antibody or a chimeric antibody.
24. (Previously Presented) The method of any of claims 21-23, wherein the antibody is an antigen-binding fragment of an antibody.
25. (Previously Presented) The method of claim 20, wherein the protein moiety is a  $\beta$ -chemokine.